

Listing of Claims:

This listing of claims will replace all prior versions and listings of claims in the application:

1-54. (Canceled)

55. (Currently amended) A solid[,] dermatological composition comprising a biologically active agent dissolved in a homogeneous carrier system, wherein the carrier system ~~includes~~ consists essentially of:

(a) 20-85% by weight of a solvent comprising (i) an unsaturated C₁₆-C₂₀-fatty acid alcohol selected from one or more of oleyl alcohol, ricinoyl alcohol, linolyl alcohol, linoleyl alcohol, eleosteryl alcohol, palmitoleyl alcohol, or arachidonyl alcohol in combination with (ii) an alkylene glycol selected from one or more of propylene glycol or dipropylene glycol, said alkylene glycol being present in the solvent in an amount of more than 12% by weight to provide for mutual dissolution of the unsaturated C₁₆-C₂₀-fatty acid alcohol and the active agent; and

~~a solvent for said active agent, comprising 20 to 85% by weight of an unsaturated fatty acid alcohol in combination with an alkylene glycol, said fatty acid alcohol being selected from oleyl alcohol, ricinoyl alcohol, linolyl alcohol and/or linoleyl alcohol, and said alkylene glycol being selected from propylene glycol and/or dipropylene glycol, said alkylene glycol being present in an amount of more than 12% by weight to provide for mutual dissolution of said active agent;~~

(b) 15-80% of a viscosity enhancing agent that is a waxy substance for imparting a solid consistency to the composition 15 to 55% by weight of a waxy substance; and

~~(c) a plasticizing agent which comprises 2 to 30% by weight of a plasticizing oil; all percentages in (a) and (b) being based on the total weight of the homogeneous carrier system.~~

56. (Currently amended) [A] The composition according to as claimed in claim 55, wherein the amount of said alkylene glycol is at least 15 % by weight.

57. (Currently amended) [A] The composition according to as claimed in claim 55, wherein the biologically active agent comprises a lipophilic compound.

58. (Currently amended) [A] The composition according to as-claimed-in claim 57, wherein the biologically active compound is selected from the group consisting of steroids, sex hormones, vitamins, biologically active lipids, fatty acids, antivirals, antibacterials, antiprotazoals, ~~and~~ antifungals, and local anesthetics.

59. (Withdrawn) A composition as claimed in claim 58, wherein the biologically active compound is selected from fluocinonide, omega-3-fatty acid and azelaic acid and salts and esters thereof.

60. (Withdrawn) A composition as claimed in claim 58, wherein the biologically active compound is clobetasol or a salt or an ester thereof.

61. (Currently amended) [A] The composition according to as-claimed-in claim 55, wherein the alkylene glycol is propylene glycol.

62. (Currently amended) [A] The composition according to as-claimed-in claim 55, wherein the solvent additionally comprises propyl myristate, palmitate, isopropylpalmitate, ~~and/or~~ stearate, ~~and/or~~ propyl ester of sorbic acid and combinations thereof.

63. (Currently amended) [A] The composition according to as-claimed-in claim 62, wherein said additional solvent is isopropylpalmitate.

64. (Currently amended) [A] The composition according to as-claimed-in claim 55, wherein the waxy substance comprises a natural ~~and/or~~ synthetic wax[:], a fat[:], a glycol ester of a C₁₈-C₃₆ fatty acid[:], or a mixture of two or more ~~such compounds~~ thereof.

65. (Currently amended) [A] The composition according to as-claimed-in claim 64, wherein the waxy substance comprises a combination of a natural or synthetic wax and one or a combination of a triglyceride or a glycol ester, ~~a triglyceride and/or a glycol ester~~.

66.-67. (Canceled)

68. (Currently amended) [A] The composition according to as-claimed-in claim 55, wherein the amount of solvent is within the range of 25-75 % by weight, and the amount of

viscosity enhancing agent is within the range of 15-55%, based on the total weight of the homogeneous carrier system.

69. (Currently amended) [A] The composition according to as-claimed-in claim 55, wherein the amount of said alkylene glycol is within the range of 12-23% by weight, based on the total weight of the homogeneous carrier system.

70. (Currently amended) [A] The composition according to as-claimed-in claim 62, wherein the weight ratio of unsaturated fatty acid : alcohol additional solvent ranges from 1:2 to 5:1.

71. (Currently amended) [A] The composition according to as-claimed-in claim 55, wherein the biologically active agent is present in a concentration of up to the solubility limit thereof in the homogeneous carrier system.

72. (Currently amended) [A] The composition according to as-claimed-in claim 55, wherein the concentration of the biologically active agent is 0.01-10%, by weight, based on the weight of the homogeneous carrier system.

73. (Currently amended) [A] The composition according to as-claimed-in claim 55, wherein said composition is a stick.

74. (Currently amended) [A] The composition according to as-claimed-in claim 55, wherein ~~said~~ the biologically active agent is a therapeutically or prophylactically active agent.

75. (Currently amended) [A] The composition according to as-claimed-in claim 74, for topical application to the skin of a mammal, ~~and possesses a~~ said composition [has] having a viscosity that is adapted for ~~such~~ said application.

76. (Currently amended) A process for the preparation of a biologically active composition ~~as-claimed-in~~ according to claim 55, comprising: dissolving the biologically active agent in said solvent therefore[,]; combining the resulting solution with a viscosity enhancing agent ~~and the plasticizing agent~~ so as to impart a solid consistency to said solution; and shaping the resulting composition into a desired form.

77. (Currently amended) A method of prophylactic or therapeutic treatment of a dermatological condition comprising: topically applying a prophylactically or therapeutically effective amount of an active agent containing the solid composition according to claim 55, wherein the active agent is an agent for treatment or prophylaxis of a dermatological condition.

78. (Currently amended) The method [of] according to claim 77, wherein the active agent is selected from the group consisting of a steroid, vitamin, biologically active lipid, fatty acid, antimicrobial, and anesthetic.

79. (Currently amended) The method [of] according to claim 77, wherein the active agent is selected from the group consisting of a corticosteroid, sex hormone, vitamin A, vitamin B2, vitamin B3, vitamin E, vitamin K, an antibiotic, an antiviral, an anti-protozoal, an antifungal, and an amide local anesthetic.

80. (Withdrawn) The method of claim 77, wherein the active agent is selected from the group consisting of clobetasol, or a salt or ester thereof and beta-methasone, or a salt or ester thereof.

81. (Withdrawn) The method of claim 80, wherein the active agent is clobetasol propionate, methasone-17-valerate, or beta-methasone dipropionate.

82. (Currently amended) [A] The composition according to as-claimed-in claim 55, wherein the biologically active agent is a lipophilic drug.

83. (Withdrawn) A composition as claimed in claim 57, wherein the biologically active compound is a lipophilic anesthetic of the amide type.

84. (Withdrawn) A composition as claimed in claim 60, wherein the biologically active compound is clobetasol propionate.

85. (Currently amended) [A] The composition according to as-claimed-in claim ~~60~~ 55, wherein the waxy substance comprises a natural ~~and/or~~ or a synthetic wax that is a monoester of a long-chain carboxylic acid with a long-chain alcohol, and the fat is a triglyceride of a C₁₈-C₃₆ fatty acid.

86.-88. (Canceled)

89. (Currently amended) [A] The composition according to as-claimed-in claim 55, wherein the amount of said alkylene glycol ranges from 15-23% by weight, based on the total weight of the homogeneous carrier system.

90. (Currently amended) [A] The composition according to as-claimed-in claim 55, wherein the amount of said alkylene glycol ranges from 12-20% by weight, based on the total weight of the homogeneous carrier system.

91. (Currently amended) [A] The composition according to as-claimed-in claim 55, wherein the amount of said alkylene glycol ranges from ranges from 15-20% by weight, based on the total weight of the homogeneous carrier system.

92. (Currently amended) [A] The composition according to as-claimed-in claim 62, wherein the weight ratio of unsaturated fatty acid alcohol : additional solvent is within the range of 1:2 to 3:1.

93. (Currently amended) [A] The composition according to as-claimed-in claim 62, wherein the weight ratio of unsaturated fatty acid alcohol : additional solvent is within the range of 1:2 to 2:1.

94. (Currently amended) [A] The composition according to as-claimed-in claim 55, wherein the concentration of the biologically active agent is 0.02-5% by weight based on the weight of the homogeneous carrier system.

95. (Withdrawn) A composition as claimed in claim 55, wherein the biologically active compound is betamethasone, or a salt or ester thereof.

96. (Withdrawn) A composition according to claim 55, wherein the biologically active compound is beta-methasone-17-valerate or beta-methasone dipropionate.

97. (Currently amended) [A] The composition according to of claim 58, wherein the steroid is a corticosteroid.

Applicants: Ake LINDAHL, *et al.*
Serial No.: 09/155,642
Filed: October 2, 1998
Page -7-

Docket No.: 28069-541 NATL
(Formerly: 402090/SkyePharma)

98. (Withdrawn) The compound of claim 58, wherein the sex hormone is selected from the group consisting of androgens, estrogens and derivatives thereof.

99. (Withdrawn) The compound of claim 58, wherein the vitamin is selected from the group consisting of vitamin A, vitamin B2, vitamin B3, vitamin E and vitamin K.